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July 1

Ar¹ and Ar² are each independently selected from the group consisting of phenyl, naphthyl, thiofuranyl, tetrahydronaphthyl, furanyl, tetrahydrofuranyl, pyridyl, quinolinyl, isoquinolinyl, tetrahydroquinolinyl, tetrahydroisoquinolinyl, cyclohexyl, cycloheptyl, and cyclopentyl;

BZ

each R¹ is independently selected from the group consisting of -H, alkyl, hydroxyalkyl, -OH, -O-alkyl, and -O-acyl;

each R² is independently selected from the group consisting of -H, alkyl, and hydroxyalkyl, or both R²s together are imino;

each R³ is independently selected from the group consisting of -H, alkyl, 2-hydroxyethyl, and alkylphenyl; and

each m is independently an integer from 0 to 5;

provided that if both R_3 's are -CH₃, then both X_m 's are not 3-F, 4-F, 3-CF₃, 4-Cl, and if both R_3 's are -CH₃ and one X_m is 4-F then the other X_m is not 4-Cl; further provided that if one R_3 is -H and the other R_3 is -CH₃ then both X_m 's are not 4-Cl, and if one R_3 is -H and the other R_3 is -CH₃ then at least one m is 1;

or a pharmaceutically acceptable salt thereof.

4. (Amended) The method of claim 3 wherein for said compound each X is independently either -F, -Cl, -OCF₃ or -CF₃;

each R¹ is -H;

each R²/is -H;

one R³ is -H, and the other R³ is either -H or -CH₃; and

each m is 1.

133 M

Atty. Dkt. No. 072827-1905

Spt.

$$(X)m$$
 Ar^1
 R^1
 R^2
 NR^3R^3
 $(X)m$
 Ar^2

133

wherein each X is independently selected from the group consisting of -Br, -Cl, -F, -I, -CF₃, alkyl, -OH, —OCF₃, -O-alkyl, and -O-acyl;

W is selected from the group consisting of -CH₂, -O-, and -S-;

Ar¹ and Ar² are each independently selected from the group consisting of phenyl, naphthyl, thiofuranyl, tetrahydronaphthyl, furanyl, tetrahydrofuranyl, pyridyl, quinolinyl, isoquinolinyl, tetrahydroguinolinyl, tetrahydroisoquinolinyl cyclohexyl, cycloheptyl, and cyclopentyl;

each R¹ is independently selected from the group consisting of -H, alkyl, hydroxyalkyl, -OH, -O-alkyl, and -O-acyl;/

each R² is independently selected from the group consisting of -H, alkyl, and hydroxyalkyl, or both R²s together are imino;

each R³ is independently selected from the group consisting of -H, alkyl, 2-hydroxyethyl, and alkylphenyl; and

m is 0 to \$;

or a pharmaceutically acceptable salt thereof.

13 4 John

$$(X)_{n}$$

$$Z$$

$$(X)_{n}$$

$$R^{1}$$

$$R^{2}$$

$$NR^{3}R^{3}$$

$$(X)_{n}$$

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wherein each X is independently selected from the group consisting of -Br, -Cl, -F, -I, -CF₃, alkyl, -OH, —OCF₃,

-O-alkyl, and -O-acyl;

each R¹ is independently selected from the group consisting of -H, alkyl, hydroxyalkyl, -O+, -O-alkyl, and -O-acyl;

each R² is independently selected from the group consisting of -H, alkyl, and hydroxyalkyl, or both R²s together are imino;

each R³ is independently selected from the group consisting of -H, alkyl, 2-hydroxyethyl, and alkylphenyl;

Z is either -CH₂CH₂-, -CH₂CH(CH₃)-, -CH=CH-, -O-CH₂-, -S-CH₂-, -CH₂-, -O-, or -S-;

each n is independently 1 to 4; or a pharmaceutically acceptable salt thereof.

135 kg

and

14. (Amended) The method of claim 13 wherein X^1 is -F, -Cl, $-OCF_3$ or $-CF_3$; and X^2 is either -F, -Cl, $-OCH_3$, $-CH_3$, $-OCF_3$ or $-CF_3$.

Sol /

p.5

$$(X)m$$
 Ar^1
 R^1
 R^2
 NR^3R^3
 $(X)m$
 Ar^2
 R^1
 R^2

wherein each X is independently selected from the group consisting of -Br, -Cl, -F, -I, -CF₃, alkyl, -OH, —OCF₃,

-O-alkyl, and -O-acyl; ; preferably, each X is independently either -F, -Cl, -OCF₃ or -CF₃; Ar¹ and Ar² are each independently selected from the group consisting of phenyl, naphthyl, thiofuranyl, tetrahydronaphthyl, furanyl, tetrahydrofuranyl, pyridyl, quinolinyl, isoquinolinyl, tetrahydroisoquinolinyl, cyclohexyl, cycloheptyl, and cyclopentyl; preferably Ar¹ and Ar² are independently naphthyl or phenyl; more preferably at least one of Ar¹ and Ar² is phenyl; and more preferably, both Ar¹ and Ar² are phenyl;

Y is either $-CH_2$ -, -O-, or -S-;

each R¹ is independently selected from the group consisting of -H, alkyl, hydroxyalkyl, -OH, -O-alkyl, and -O-acyl; preferably, each R¹ is -H;

each R² is independently selected from the group consisting of -H, alkyl, and hydroxyalkyl, or both R²s together are imino; preferably each R² is -H;

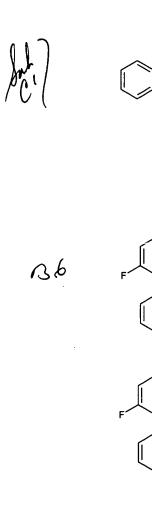
each R³ is independently selected from the group consisting of -H, alkyl, 2-hydroxyethyl, and alkylphenyl; preferably, each R³ is independently either -H or -CH₃; more preferably one R³ is

-H, and the other R³ is either -H or -CH; and

each m is independently an integer from 0 to 5; and preferably, each m is independently 0

or 1.

S Col



-7-

Please add the following new claims:

n 7

19. (New) The method of claim 3 wherein said compound has the chemical structure:

$$X^{1} \stackrel{\stackrel{}{\parallel}}{\longrightarrow} R^{2} \stackrel{\stackrel{}{\parallel}}{\longrightarrow} NR^{3}R^{3}$$

$$X^{2} \stackrel{\stackrel{}{\parallel}}{\longrightarrow} R^{2}$$

wherein

X¹ is either -Br, -Cl, -F, -I, -CF₃/alkyl, -OH, —OCF₃, -O-alkyl, or -O-acyl;

 X^2 is either -Br, -Cl, -F, -I, -C $\sqrt{3}$, alkyl, -OH, —OCF₃, -O-alkyl, or -O-acyl;

each R¹ is independently selected from the group consisting of -H, alkyl, hydroxyalkyl, -O+, -O-alkyl, and -O-acyl;

each R² is independently selected from the group consisting of -H, alkyl, and hydroxyalkyl, or both R²s together are imino

each R³ is independently selected from the group consisting of -H, alkyl, 2-hydroxyethyl, and alkylphenyl;

or a pharmaceutically acceptable salt thereof.

20. The method of claim 19, wherein

each X is independently either -F, -Cl, -OCF₃ or -CF₃;

each R' is -H;

each/R² is -H; and

one \mathbb{R}^3 is -H, and the other \mathbb{R}^3 is either -H or -CH₃.